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DOCKET NO.: 9491-057-27 CONT

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Re: Serial No.: 09/995,693
Applicant(s): RALPH H. SCHWALL, ET AL.
Filing Date: NOVEMBER 29, 2001
For: HEPATOCYTE GROWTH FACTOR RECEPTOR ANTAGONISTS
AND USES THEREOF
Group Art Unit: 1642
Examiner: UNASSIGNED

SIR:

Attached hereto for filing are the following papers:

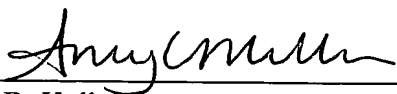
NOTICE TO COMPLY (RETURN COPY)
RESPONSE TO NOTICE TO COMPLY AND SECOND SUPPLEMENTAL
PRELIMINARY AMENDMENT INCLUDING ATTACHMENT
REQUEST FOR PREPARATION OF COMPUTER-READABLE SEQUENCE LISTING
STATEMENT UNDER 37 C.F.R. §1.821-1.825 (1 PAGE, EXECUTED)
SUBSTITUTE SEQUENCE LISTING (PAPER COPY, 7 PAGES)

Our check in the amount of \$ -0- is attached covering any required fees. In the event any variance exists between the amount enclosed and the Patent Office charges for filing the above-noted documents, including any fees required under 37 C.F.R. 1.136 for any necessary extension of time to make the filing of the attached documents timely, please charge or credit the difference to Deposit Account No. 50-1442. Further, if these papers are not considered timely filed, then a request is hereby made under 37 C.F.R. 1.136 for the necessary extension of time. A duplicate copy of this sheet is enclosed.

Respectfully submitted,

PIPER MARBURY RUDNICK & WOLFE LLP

Date: 2-19-02



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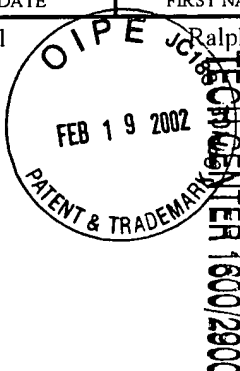


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APPLICATION NUMBER	FILING/RECEIPT DATE	FIRST NAMED APPLICANT	ATTORNEY DOCKET NUMBER
09/995,693	11/29/2001	Ralph H. Schwall	9491-057-27 CONT

Supervisor, Patent Prosecution Services
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CONFIRMATION NO. 3638

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Applicant is given **TWO MONTHS FROM THE DATE OF THIS NOTICE** within which to file the items indicated below to avoid abandonment. Extensions of time may be obtained under the provisions of 37 CFR 1.136(a).

- A request to transfer the computer readable form from another application on file at the U.S. Patent and Trademark Office has been submitted as permitted by 37 C.F.R. 1.821(e). However, the request cannot be complied with since there is no compliant CRF present at the United States Patent and Trademark Office. Applicant must provide an initial computer readable form (CRF) copy of the "Sequence Listing" and a statement that the content of the sequence listing information recorded in computer readable form is identical to the written (on paper or compact disc) sequence listing and, where applicable, includes no new matter, as required by 37 CFR 1.821(e), 1.821(f), 1.821(g), 1.825(b), or 1.825(d).

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